

Amendments to the Claims:

Please amend claim 9 and cancel claims 35, 40-50. This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

1. (Withdrawn) A method for identifying a compound capable of interfering with binding of a SAK polypeptide or fragment thereof, the method comprising the steps of:
 - (i) combining a SAK polypeptide or fragment thereof with a Chk2 polypeptide and the compound, wherein the SAK polypeptide or fragment thereof has kinase activity and is encoded by a nucleic acid that hybridizes under stringent conditions to a nucleic acid encoding a polypeptide having an amino acid sequence of SEQ ID NO:2; and
 - (ii) determining the binding of the SAK polypeptide or fragment thereof to Chk2.
2. (Withdrawn) The method of claim 1, wherein the SAK polypeptide or fragment thereof and the Chk2 polypeptide are combined first.
3. (Withdrawn) The method of claim 1, wherein the binding of the SAK polypeptide or fragment thereof to Chk2 is determined in vitro.
4. (Withdrawn) The method of claim 1, wherein the SAK polypeptide or fragment thereof and the Chk2 polypeptide are expressed in a cell.
5. (Withdrawn) The method of claim 4, wherein the cell is a yeast or a mammalian cell.
6. (Withdrawn) The method of claim 5, wherein the SAK polypeptide or fragment thereof is fused to a heterologous polypeptide.
7. (Withdrawn) The method of claim 1, wherein the binding of the SAK polypeptide or fragment thereof to Chk2 is determined by measuring reporter gene expression.

8. (Withdrawn) The method of claim 1, wherein the binding of the SAK polypeptide or fragment thereof to Chk2 is determined by measuring SAK kinase activity.

9. (Currently Amended) A method for identifying a compound that modulates cellular proliferation, the method comprising the steps of:

(i) contacting the compound with a SAK polypeptide, the polypeptide encoded by a nucleic acid that ~~hybridizes under stringent conditions to a nucleic acid encoding a polypeptide having an amino acid sequence of SEQ ID NO:2 that encodes a SAK polypeptide having at least 95% sequence identity to SEQ ID NO:2, the polypeptide having serine/threonine kinase activity;~~ and

(ii) determining the functional effect of the compound upon the SAK polypeptide; and

(iii) identifying the compound based on step (ii).

10. (Original) The method of claim 9, wherein the functional effect is measured in vitro.

11. (Original) The method of claim 10, wherein the functional effect is a physical effect.

12. (Withdrawn) The method of claim 11, wherein the physical effect is determined by measuring ligand or substrate binding to the polypeptide.

13. (Withdrawn) The method of claim 10, wherein the functional effect is a chemical effect.

14. (Withdrawn) The method of claim 13, wherein the chemical effect is determined by measuring kinase activity of the SAK polypeptide.

15. (Original) The method of claim 9, wherein the polypeptide is expressed in a eukaryotic host cell.

16. (Original) The method of claim 15, wherein the functional effect is a physical effect.

17. (Withdrawn) The method of claim 16, wherein the physical effect is determined by measuring ligand or substrate binding to the polypeptide.

18. (Original) The method of claim 15, wherein the functional effect is a chemical or phenotypic effect.

19. (Withdrawn) The method of claim 18, wherein the chemical or phenotypic effect is determined by measuring kinase activity of the SAK polypeptide.

20. (Original) The method of claim 18, wherein the chemical or phenotypic effect is determined by measuring cellular proliferation.

21. (Original) The method of claim 20, wherein the cellular proliferation is measured by assaying for DNA synthesis or fluorescent marker dilution.

22. (Original) The method of claim 21, wherein DNA synthesis is measured by 3H thymidine incorporation, BrdU incorporation, or Hoescht staining.

23. (Original) The method of claim 21, wherein the fluorescent marker is selected from the group consisting of a cell tracker dye or green fluorescent protein.

24. (Original) The method of claim 9, wherein modulation is inhibition of cellular proliferation.

25. (Original) The method of claim 9, wherein modulation is inhibition of cancer cell proliferation.

26. (Original) The method of claim 15, wherein the host cell is a cancer cell.

27. (Original) The method of claim 26, wherein the cancer cell is a breast, prostate, colon, or lung cancer cell.

28. (Original) The method of claim 26, wherein the cancer cell is a transformed cell line.

29. (Original) The method of claim 28, wherein the transformed cell line is PC3, H1299, MDA-MB-231, MCF7, A549, or HeLa.

30. (Original) The method of claim 26, wherein the cancer cell is p53 null or mutant.

31. (Original) The method of claim 26, wherein the cancer cell is p53 wild-type.

32. (Original) The method of claim 9, wherein the polypeptide is recombinant.

33. (Original) The method of claim 9, wherein the polypeptide is encoded by a nucleic acid comprising a sequence of SEQ ID NO:1.

34. (Original) The method of claim 9, wherein the compound is an antibody.

35. (canceled)

36. (Original) The method of claim 9, wherein the compound is a small organic molecule.

37. (Original) The method of claim 9, wherein the compound is a peptide.

38. (Original) The method of claim 37, wherein the peptide is circular.

39. (Withdrawn) A method for identifying a compound that modulates cellular proliferation or chemosensitivity, the method comprising the steps of:

(i) contacting the compound with an SAK polypeptide or a fragment thereof, the SAK polypeptide or fragment thereof encoded by a nucleic acid that hybridizes under stringent conditions to a nucleic acid encoded by a polypeptide comprising an amino acid sequence of SEQ ID NO:2;

(ii) determining the physical effect of the compound upon the SAK polypeptide; and

(iii) determining the chemical or phenotypic effect of the compound upon a cell comprising an SAK polypeptide or fragment thereof, thereby identifying a compound that modulates cellular proliferation or chemosensitivity.

40-50. (canceled)